## Introduction

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**Dotplot** is a tool for comparing the sequences of two genomes. **Dotplot** employs the program **Mummer** to generate dotplot diagrams between two genomes. It uses input DNA sequences directly for comparing genomes with similar sequences (<u>NUCmer</u>). It uses the six frame amino acid translation of the DNA input sequences (<u>PROmer</u>) for comparing genomes with dissimilar sequences (because the DNA sequence is not as highly conserved as the amino acid translation).

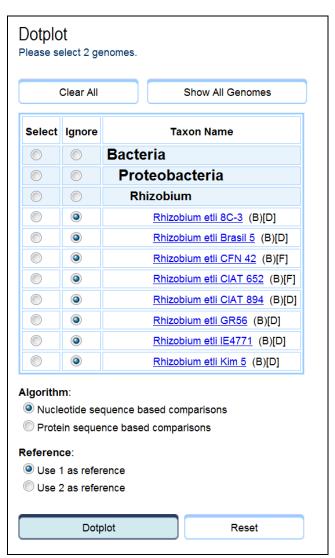


Figure 1. Select 2 genomes for Dot Plot

**Dotplot** can be applied on two genomes that are selected from the list of all available or preselected genomes (Figure 1). Select which **algorithm** to use (nucleotide based comparison for genomes with similar nucleotide sequences or protein base comparison for genomes with dissimilar nucleotide sequences), and select the genome that will be used as **reference**.

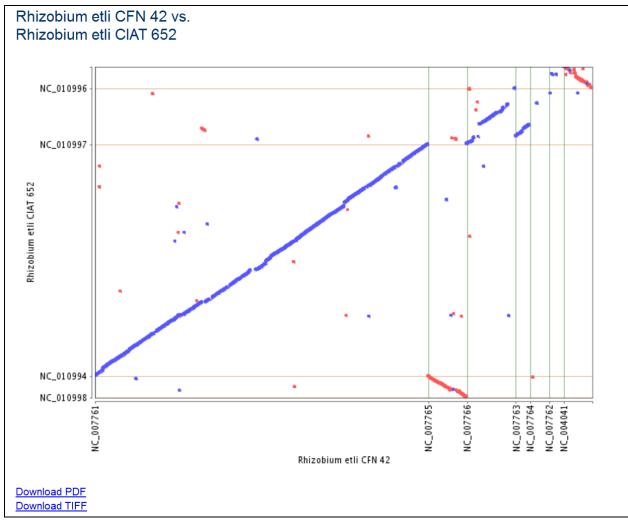


Figure 2. Dot Plot – fplot (blue), rplot (red)

The dotplot (Figure 2) consists of **blue** points for regions of similarity found on parallel strands (fplot) and **red** points for regions of similarity found on antiparallel strands (rplot). The tooltip for each point shows which plot it is and the coordinates and scaffolds of the alignment. Clicking on a point brings up the chromosomal neighborhood of the alignment of the two genomes (Figure 3). The plot can be saved either as PDF or as TIFF.

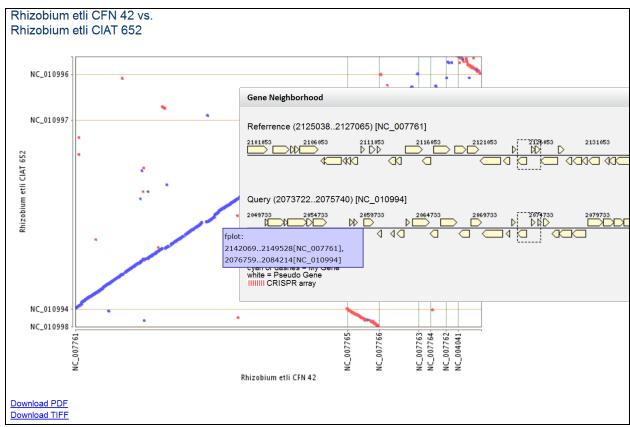


Figure 3. Dot Plot – tooltips, links to Gene Neighborhood of aligned region

## Scaffold Selection MG 3.3

When the number of scaffolds for the 2 genomes selected for Dot Plot is too high, the computation of the Dot Plot may time out. In such a scenario, the user is offered a choice to select a subset of scaffolds or to continue with the calculation using all scaffolds (Figure 5). The current cutoff is when the number of scaffold is >150 for both genomes or >300 for one of the genomes

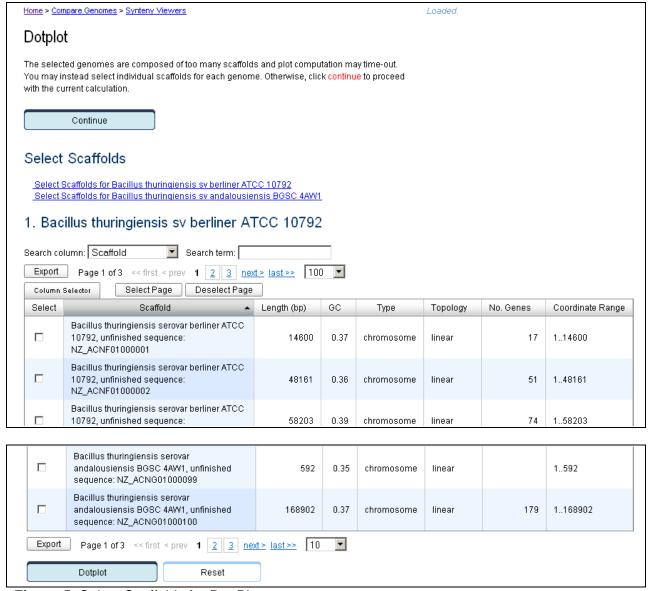


Figure 5. Select Scaffolds for Dot Plot